

PURELL ANTIMICROBIAL FOAMING HAND SP WITH PCMX- chloroxylenol liquid
GOJO Industries, Inc.

PURELL HEALTHY SOAP 0.5% PCMX Antimicrobial Foam

Active Ingredient

Chloroxylenol 0.5%

Purpose

Microbial

Use

- Handwash to help decrease bacteria on the skin.
- Recommended for repeated use.

Warnings

When using this product do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash appears and lasts.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Wet hands
- Apply a small amount of soap and work into a rich lather
- Rinse thoroughly with potable water
- Dry hands completely.

Inactive Ingredients

Water (Aqua), Alcohol, Lauric Acid, Ethanolamine, Dipropylene Glycol, Lactic Acid, Poloxamer 124, Isopropyl Alcohol, Sodium Metabisulfite, Tetrasodium EDTA, Fragrance (Parfum), Methylparaben, Propylparaben



ANTIMICROBIAL
**HAND
SOAP**

99.9%
KILLS 99.9%
OF GERMS*


HYPOALLERGENIC


GENTLE
ON SKIN



0 73852 30785 6



**Antimicrobial Foaming
Hand Soap with PCMX**
Jabón de Manos Espumoso
Antimicrobiano con PCMX

Drug Facts	
Active Ingredient	Purpose
Chloroxyleneol 0.5%	Antimicrobial
Use • Handwash to help decrease bacteria on the skin. • Recommended for repeated use.	
Warnings For external use only	
When using this product do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.	
Stop use and ask a doctor if irritation or rash appears and lasts.	
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.	
Directions • Wet hands • Apply a small amount of soap and work into a rich lather. • Rinse thoroughly with potable water. • Dry hands completely.	
Inactive ingredients: Water (Aqua), Alcohol, Lauric Acid, Ethanolamine, Dipropylene Glycol, Lactic Acid, Poloxamer 124, Isopropyl Alcohol, Sodium Metabisulfite, Tetrasodium EDTA, Fragrance (Parfum), Methylparaben, Propylparaben.	

Datos Farmacológicos

Ingrediente activo	Propósito
Cloroxilenol 0,5%	Antimicrobiano
Uso • Lavado de manos enjabonado para disminuir la cantidad de bacterias en la piel. • Recomendado para uso repetido.	
Advertencias Solo para uso externo	
Al utilizar este producto , evitar el contacto con los ojos o con la zona alrededor de los ojos. En caso de contacto, enjuagar completamente los ojos con agua.	
Dejar de usar el producto y consultar a un médico si aparece y persiste una irritación o erupción cutánea.	
Mantener fuera del alcance de los niños. En caso de ingestión, se recomienda acudir a un médico o ponerse en contacto con un centro para el control de tóxicos.	
Modo de uso • Mojar las manos • Aplicar una pequeña cantidad de jabón y frotar las manos hasta producir una espuma abundante. • Enjuagar bien con agua potable. • Secarse las manos completamente.	
Ingredientes inactivos: Agua, Alcohol etílico, Ácido láurico, Etanolamina, Cloropiridol, Ácido láctico, 4-cloxiámero 124, Alcohol isopropílico, Metabisulfito de sodio, EDTA tetrasódico, Fragancia, Metilparabeno, Propilparabén.	

1.2 L (40.5 US/ÉU FL OZ)



Reorder No.
Código N° 8384 DSP-OH-36



PURELL ANTIMICROBIAL FOAMING HAND SP WITH PCMX

chloroxylenol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:21749-722
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	0.5 mg in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	
LAURIC ACID (UNII: 1160N9NU9U)	
MONOETHANOLAMINE (UNII: 5KV86114PT)	
DIPROPYLENE GLYCOL (UNII: E107L85C40)	
LACTIC ACID (UNII: 33X04XA5AT)	
POLOXAMER 124 (UNII: 1S66E28KXA)	

ISOPROPYL ALCOHOL (UNII: ND2M416302)				
SODIUM METABISULFITE (UNII: 4VON5FNS3C)				
EDETATE SODIUM (UNII: MP1J8420LU)				
METHYLPARABEN (UNII: A2I8C7HI9T)				
PROPYLPARABEN (UNII: Z8IX2SC1OH)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:21749-722-89	1200 mL in 1 PACKAGE; Type 0: Not a Combination Product	12/01/2025	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug		505G(a)(3)	12/01/2025	

Labeler - GOJO Industries, Inc. (004162038)

Establishment

Name	Address	ID/FEI	Business Operations
GOJO Industries, Inc.		036424534	manufacture(21749-722) , pack(21749-722) , label(21749-722)

Revised: 11/2025

GOJO Industries, Inc.